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Amendments To The Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1 (currently amended). A composition comprising [[a]] first oligomeric compound and a second oligomeric compound compounds wherein:

at least a portion 18 nucleosides of the first oligomeric compound is are capable of hybridizing with at least a portion of the second oligomeric compound;

at least a portion of the first oligomeric compound is complementary to and capable of hybridizing to a target nucleic acid; and

at least one of the first and the second oligomeric compounds comprises at least one sugar modified nucleoside having enhanced or decreased affinity for the complementary nucleoside in the composition or between the first oligomeric compound and a nucleic acid target relative to an unmodified β-D-ribonucleoside; or and

one the other of the first and the second oligomeric compounds comprises at least one base modified nucleoside having enhanced affinity for the complementary nucleoside in the composition or between the first oligomeric compound and a nucleic acid target and one of the first and the second oligomeric compounds comprises at least one modified nucleoside having decreased affinity for the complementary nucleoside in the composition or between the first oligomeric compound and a nucleic acid target relative to an unmodified \(\mathcal{B} \- \mathcal{D} \- \text{ribonucleoside} \).

2 (currently amended). The composition of claim 1 wherein the first oligomeric compound comprises <u>said</u> at least one <u>sugar</u> modified nucleoside for the complementary nucleoside in the composition or between the first oligomeric compound and a nucleic acid target and either the first oligomeric compound or second oligomeric compound comprises at least one modified nucleoside having a decreased affinity for the complementary nucleoside in the composition or between the first oligomeric compound and a nucleic acid target.

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3 (currently amended). The composition of claim 1 wherein the first oligomeric compound comprises at least one modified nucleoside having a decreased affinity for the complementary nucleoside in the composition or between the first oligomeric compound and a nucleic acid target, and the second oligomeric compound comprises said at least one sugar modified nucleotide having an enhanced affinity for the complementary nucleotide in the first oligomeric compound compared to the affinity of an unmodified nucleotide.

4 (currently amended). The composition of claim ± 2 wherein the second oligomeric compound comprises at least one <u>base</u> modified <u>nucleotide</u> <u>nucleoside</u> having an enhanced <u>decreased</u> affinity for the complementary nucleotide in the first oligomeric compound compared to the affinity of an unmodified nucleotide, and wherein the second oligomeric compound also comprises at least one modified nucleotide having a decreased affinity for the complementary nucleotide in the first oligomeric compound compared to the affinity of an unmodified nucleotide.

5 (currently amended). The composition of claim 1 wherein the <u>second oligomeric</u>

<u>compound comprises</u> at least one <u>base</u> modified <u>nucleotide</u> <u>nucleoside</u> that <u>comprises an having</u>

enhanced affinity is a nucleotide comprising a nucleotide base modification.

6 (currently amended). The composition of claim 5 wherein the <u>nucleotide nucleoside</u> base modification comprises a pyrimidine <u>nucleotide nucleoside</u> comprising a modification <u>modified</u> at the 2, 4, 5 or 6 position of the pyrimidine nucleotide.

7 (currently amended). The composition of claim 6 wherein the pyrimidine nucleotide nucleoside comprises a modification at the 2 or 5 position of the pyrimidine nucleotide.

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8 (currently amended). The composition of claim 6 wherein the <u>nucleotide nucleoside</u> base modification comprises a 2-thio U <u>nucleotide substitution for U nucleotide</u> or 2-thio C <u>nucleotide</u> substitution for a C nucleotide.

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9 (currently amended). The composition of claim 6 wherein the <u>nucleotide nucleoside</u> base modification comprises a 5-alkyl, 5-alkenyl, or 5-alkynyl U <u>substitution for a U nucleotide</u> or 5-alkyl, 5-alkenyl, or 5-alkynyl C <u>substitution for a C nucleotide</u>.

10 (currently amended). The composition of claim 6 wherein the <u>nucleotide</u> base modification comprises a 5-methyl U, 5-methyl C, 5-propynyl U, or 5-propynyl C <u>nucleotide</u>.

11 (currently amended). The composition of claim 5 wherein the <u>nucleotide nucleoside</u> base modification comprises a <u>pyrimidine having a modification</u>, wherein the <u>pyrimidine nucleotide is incorporated as one ring of a multiple ring heterocycle</u>, wherein the multiple ring heterocycle further comprises a phenoxazine moiety, and wherein the multiple ring heterocycle comprises the formula:

wherein:

 R_{11} is $(CH_3)_2N-(CH_2)_2-O-$; $H_2N-(CH_2)_3-$; $Ph-CH_2-O-C(=O)-N(H)-(CH_2)_3-$; H_2N- ; fluorenyl-CH₂-O-C(=O)-N(H)-(CH₂)₃-; $Ph-CH_2-O-C(=O)-N(H)-(CH_2)_3-$; $Ph-CH_2-O-C(=O)-N(H)-(CH_2)_2-O-$; $Ph-CH_2-O-C(=O)-N(H)-(CH_2)_3-O-$; $Ph-CH_2-O-C(EO)-N(H)-(EH_2)_3-O-$;

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fluorenyl-CH₂-O-C(=O)-N(H)-(CH₂)₂-O-; fluorenyl-CH₂-O-C(=O)-N(H)-(CH₂)₃-O-; H₂N-(CH₂)₂-O-CH₂-; N₃-(CH₂)₂-O-CH₂-; H₂N-(CH₂)₂-O-, or NH₂C(=NH)NH-.

12-13. (cancelled).

14 (original). The composition of claim 5 wherein the <u>nucleotide</u> <u>nucleoside</u> base modification comprises a purine <u>nucleotide</u> <u>nucleoside</u> <u>comprising a modification</u> <u>modified</u> at the 1, 2, 3, 6, 7 or 8 position of the purine nucleotide.

15 (currently amended). The composition of claim 14 wherein the nucleotide nucleoside base modification comprises a purine nucleotide comprising a modification modified at the 2, 6 or 7 positions of the purine nucleotide position.

16. (cancelled).

17 (currently amended). The composition of claim 14 wherein the nucleotide nucleoside base modification comprises a 2,6-diamino purine substitution for an A nucleotide.

18 (canceled).

19 (currently amended). The composition of claim 18 1 wherein the nucleotide sugar modification comprises a 2'-substituent group selected from 2'-F, 2'-MOE (2-O-(CH₂)₂-OCH₃), 2'-O-methyl, 2'-O-alkyl, 2'-O-alkynyl, 2'-S-alkyl, 2'-S-alkyl, 2'-S-alkyl, 2'-S-alkynyl, 2'-amino, 2'-azido, or 2'-allyl.

20-23 (canceled).

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24 (currently amended). The composition of claim 23 1 wherein the nucleotide base modified nucleoside modification comprises an is inosine nucleotide or a purine ribofuranosyl nucleotide.

25-29 (canceled).

30. (currently amended). The composition of claim $5\ \underline{1}$ wherein the nucleotide base sugar modification comprises a 2'-substituent group which is, independently, F, -O-CH₂CH₂-O-CH₃, -O-C₁-C₁₂ alkyl, -O-CH₂-CH₂-CH₂-NH₂, -O-(CH₂)₂-O-N(R₁)₂, -O-CH₂C(=O)-N(R₁)₂, -O-(CH₂)₂-O-(CH₂)₂-O-(CH₂)₂-N(R₁)₂, -O-CH₂-CH₂-CH₂-NHR₁, -O-CF₃, -N₃, -O-CH₂-CH=CH₂, -NHCOR₁, -NH₂, -NHR₁, -N(R₁)₂, -SH, -SR₁, -N(H)OH, -N(H)OR₁, -N(R₁)OH, -N(R₁)OR₁ or -O-CH₂-N(H)-C(=NR₁)(N(R₁)₂);

wherein each R_1 is, independently, H, C_1 - C_{12} alkyl, a protecting group, or substituted or unsubstituted C_1 - C_{12} alkyl, C_2 - C_{12} alkenyl, or C_2 - C_{12} alkynyl, wherein the substituent groups are halogen, hydroxyl, amino, azido, cyano, haloalkyl, alkenyl, alkoxy, thioalkoxy, haloalkoxy, or aryl.

31 (original). The composition of claim 30 wherein each of the 2'-substituent groups is, independently, -F, -O-CH₃, -O-CH₂CH₂-O-CH₃, -O-CH₂-CH=CH₂, -O-CF₃, N₃, NH₂, NHOH, -O-(CH₂)₂-O-N(R₁)₂, -O-CH₂C(O)-N(R₁)₂, -O-CH₂-CH₂-CH₂-NH₂, -O-(CH₂)₂-O-(CH₂)₂-N(R₁)₂ or -O-CH₂-N(H)-C(=NR₁)(N(R₁)₂);

wherein each R_1 is, independently, H, C_1 - C_{12} alkyl, a protecting group, or substituted or unsubstituted C_1 - C_{12} alkyl, C_2 - C_{12} alkenyl, or C_2 - C_{12} alkynyl, wherein the substituent groups are halogen, hydroxyl, amino, azido, cyano, haloalkyl, alkenyl, alkoxy, thioalkoxy, haloalkoxy, or aryl.

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32 (original). The composition of claim 31 wherein each of the 2'-substituent groups is, independently, -F, -O-CH₂CH₂-O-CH₃, -O-CH₃, -O-CH₂-CH=CH₂, -O-CF₃ or -O-CH₂-CH-CH₂-NH(R_j) where R_j is H or C₁-C₁₀ alkyl.

33 (original). The composition of claim 32 wherein each of the 2'-substituent groups is, independently, F, -O-CH₃, -O-CF₃, or -O-CH₂CH₂-O-CH₃.

34. (currently amended). The composition of claim 5 1 wherein at least one <u>sugar</u> modified nucleotide <u>nucleoside</u> base is a locked nucleic acid (LNA).

35-37 (canceled).

38 (currently amended). The composition of claim 1 wherein each of the first and second oligomeric compounds comprises from about 12 to about 30 nucleobases nucleosides.

39 (canceled).

40 (currently amended). The composition of claim 1 wherein each of the first and second oligomeric compounds comprises from about 19 to about 23 nucleosides.

41-56 (canceled).

57 (new). The composition of claim 1 wherein each of said sugar modified nucleosides is a 2'-F modified nucleosides nucleoside.

58 (new). The composition of claim 57 wherein one of the first and the second oligomeric compounds comprises a continuous sequence of 2'-F modified nucleosides.

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59 (new). The composition of claim 58 wherein the other of the first and the second oligomeric compounds comprises at least one base modified nucleoside having said decreased affinity.

60 (new). The composition of claim 59 wherein each of the base modified nucleosides having said decreased affinity is an inosine modified nucleoside.